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ABSTRACT

Novel drug complexes comprising a polypeptide carrier moiety comprising glutamic acid and at least one of the group consisting of aspartic acid, alanine, asparagine, glutamine, glycine, and any combinations thereof, are disclosed. The drug moiety is a therapeutic metal selected from the group consisting of platinum, iron, gadolinium, rhenium, manganese, cobolt, indium, gallium or Methods for making said complexes, rhodium. compositions comprising said complexes, methods for making saiduch compositions, and methods for treating a patient comprising use of said complexes and/or compositions are further disclosed.

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Mol Percent Report

Sample IO: 01402018 (initiated 1/15/99 7:03am) BASELINE CORRECTED Turntable Position: e c Sampling Interval: 1.0 sec Data Start 3.00 min Samples In Run : 72 Data Duration : 16.00 min Operator ID Peak Ht Thrashold: 3000 uAU Int. Std. Amt : 250 pmol Calibration File : 14JANCAL (initiated 1/15/99 9:44am) Reference Time : 0.00 min (No ISTD Peak Specified)

Reference Offset 1: -0.02 min Reference Offset Z: 0.00 min

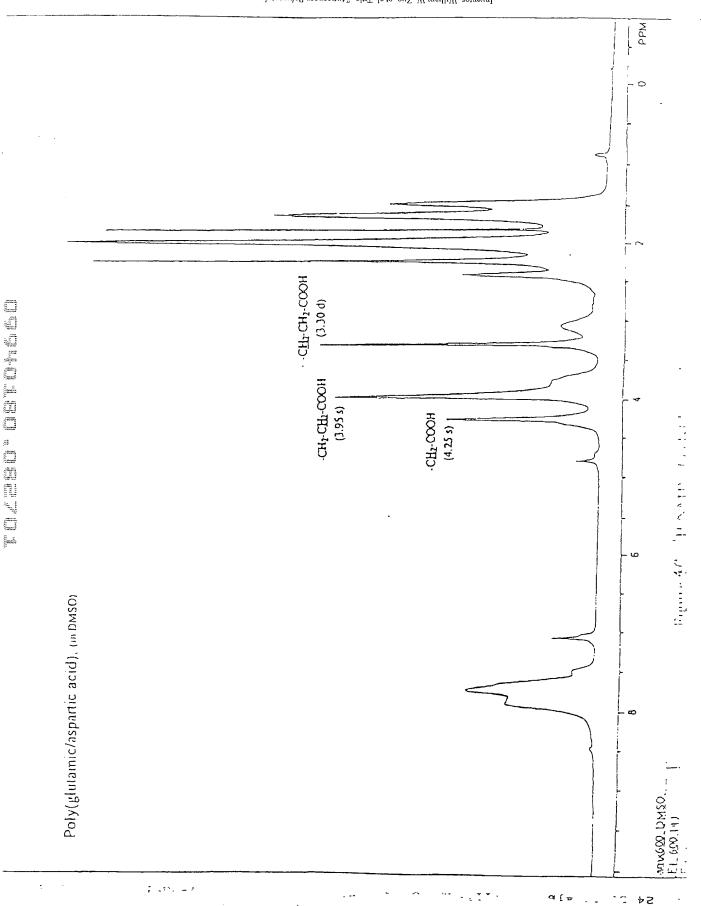
Integration Interval: 0.0 to 16.0 min

PEAK ID	RET. TIME	BY BY		PMOL correc.	MOL	
	Miπ	HEIGHT		INT STO	X.	
ASX	2.30	562.99		0.00	29.02	
GLX	2.70	1377.05		0.00	70.98	
TOTAL PHOLS	RECOVERED	1940.04				
Minimum Peak	Threshold:	3000 uAU	(5	3 peaks below	threshold)
			(1	8 peaks found		}
			(2 peaks match	ed)

Figure 1B.

Amino acid analysis of the polypeptide

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11:0%

Pt(II) Complex

Pt(IV) Complex

Figure 2.

Structures of platinum(II) and platinum(IV)-poly(dipeptide) complexes



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LABORATORY REPORT

SAMPLE ID	LABID	ANALYSIS	RESULT(S)	
DACH	1-5746	Plannum	44 60	C y
PPAP	1-5747	Platinum	16.11	v o
PDDP	1-5748	Platinum	1764	٠,

Figure ->

Elemental analysis of platinum-poly(dipeptide) complexes for Pt(II)(PDDP), Pt(IV)(PPAP) and cis-1,2-DACH-Pt SO₄(DACH)

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Inventor William W Zuo, et al Title "Anticancer Polypeptide Metal Complexes and Compositions, Methods of Making, and Methods of Using Same"

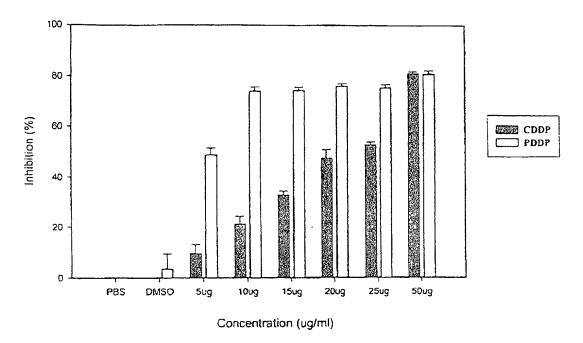
Agent Mary A Gilbreth, Ph D Reg No 45,775

Atty Docket 27300/03

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Effect of PDDP & CDDP on Inhibition of Human Sarcoma Cells (HT1080) at 48 Hours

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Effect of PDDP & CDDP on Inhibition of Human Sarcoma Cells (HT1080) at 72 Hours

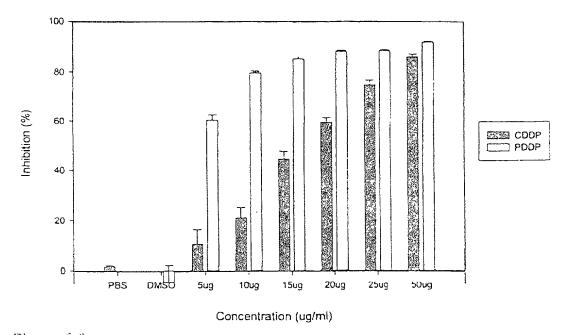
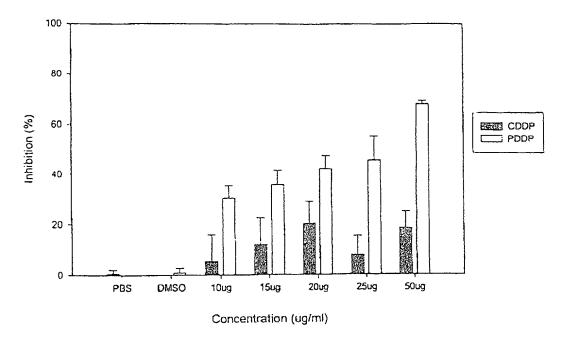


Figure 4A

In vitro cell culture assay of cisplatin(CDDP) and poly(glutamate/aspartate) acid-1,2-DACH-Pt(II) complex(PDDP) in human sarcoma

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Effect of PDDP & CDDP on Inhibition of Human Prostate Cancer Cells (A10) at 48 Hours



Effect of PDDP &CDDP on Inhibition of Human Prostate Cancer Cells (A10) at 96 Hours

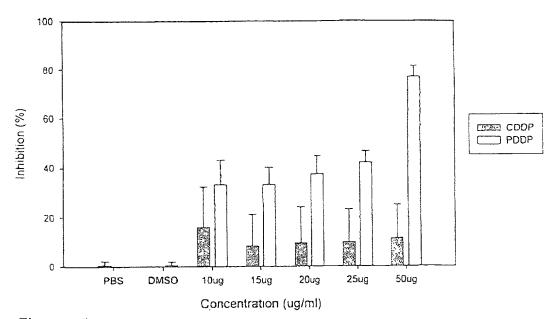
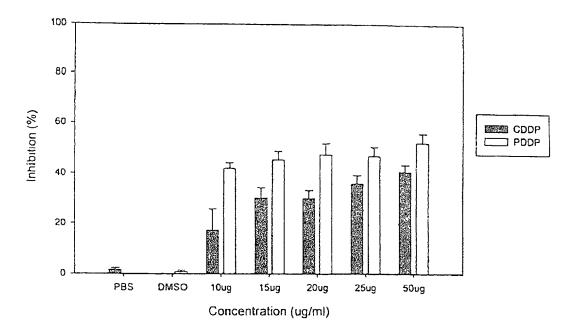


Figure 4-6 In vitro cell culture assay of cisplatin(CDDP) and poly(glutamate/aspartate) acid-1,2-DACH-Pt(II) complex(PDDP) in human prostate cancer

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Effect of PDDP & CDDP on Inhibition of Human Prostate Cancer Cells (PC3) at 48 Hours



Effect of PDDP & CDDP on Inhibition of Human Prostate Cancer Cells (PC3) at 96 Hours

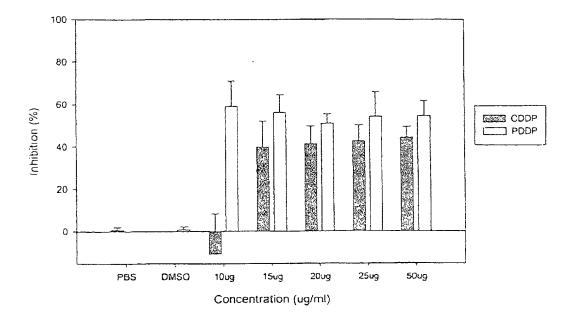
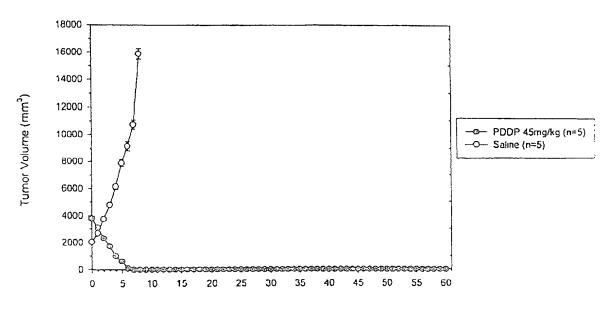


Figure 50.

In vitro cell culture assay of cisplatin(CDDP) and poly(glutamate/aspartate) acid-1,2-DACH-Pt(II) complex(PDDP) in human prostate cancer

Anti-tumor Activity of PDDP Against Rats Bearing Breast Tumors (13762)



Days after Treatment (single injection, i.v.)

Figure 5
In vivo anti-tumor activity of an inventive poly(glutamate/aspartate) acid-1,2-DACH-Pt(II) complex (PDDP) compared to control against rats bearing breast tumors.

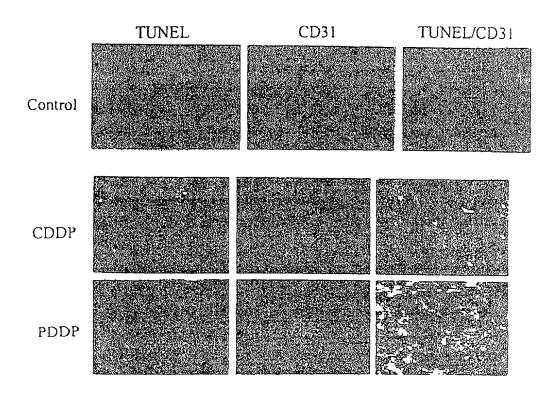
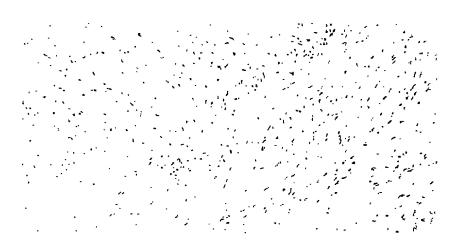
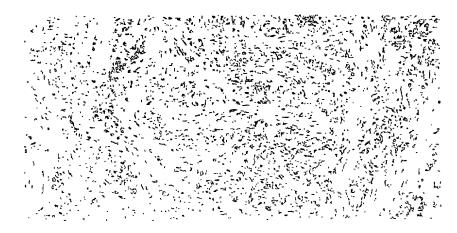


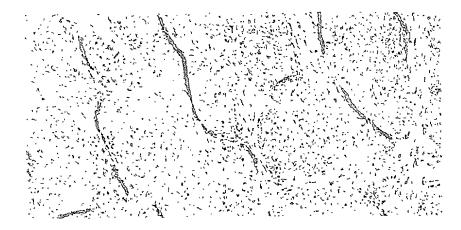
Figure $\mathfrak S$ The specific cellular target expression changes at 48 hours post treatment of poly(glutamate/aspartate) acid-1,2-DACH-Pt(II) complex (PDDP), cisplatin (CDDP) and saline (control)



Histopathological slide of the breast tumor (13762) at 48 hours post treatment of PDDP



Histopathological slide of the breast tumor (13762) at 48 hours post treatment of CDDP



Histopathological slide of the breast tumor (13762) at 48 hours post treatment of Salme Figure \mp

The histopathological changes of breast tumors (13762) at 48 hours post treatment of poly(glutaniate/aspartate) acid-1.2-DACH-Pt(II) complex (PDDP), cisplatin (CDDP) and Saline